



Atomic Structure and Properties Unit 1 ↓	Molecular & Ionic Compound Structure and Properties Unit 2 ↓	Intermolecular Forces & Properties Unit 3 ↓	Chemical Reactions Unit 4 ↓	Kinetics Unit 5 ↓
<ul style="list-style-type: none"> <u>Conversions</u> - Avogadro's number, molar mass, and mole ratios <u>Empirical+Molecular Formula</u> - These are the simplest whole # ratio of atoms for a compound and the chemical formula for a compound, respectively. <u>Mass Spectroscopy</u> - Mass to charge ratio of compounds. <u>Electron Configurations</u> - Electrons fill the lowest energy level orbital first, no two e⁻ can have the same spin, and e⁻ occupy separate subshells before sharing one. <u>Photoelectron Spectroscopy</u> - Measures the amount of energy electrons release. <u>Periodic trends</u> - recognizing them and explaining them <u>Mixtures</u> - Homogeneous (pure) and heterogeneous 	<ul style="list-style-type: none"> <u>Ionic Bonds</u> - between metal and nonmetals, e⁻ are transferred. <u>Covalent Bonds</u> - between nonmetals, e⁻ are shared. <u>Lattice Energy</u> - energy of ionic bonds. <u>Metallic Bonds</u> - The sharing of free e⁻ between metal atoms. <u>Alloys</u> - Compounds of different metals <u>Lewis Structures and VSEPR</u> - Bonding diagrams and geometric, 3-D shapes of compounds. <u>Hybridization</u> - atomic orbitals fuse to form new orbitals <u>Formal Charge</u> - Charge of an element in a molecule. <u>Resonance</u> - Molecules bonding structure is a combination of other possible structures. <u>Coulomb's Law</u> - shorter distances + higher charges = strongest attractions 	<p>Strongest to Weakest IMFs:</p> <ul style="list-style-type: none"> <u>Ion-Dipole</u> - ionic compounds + liquid <u>H-Bonds</u> - fluorine, oxygen, nitrogen <u>Dipole-Dipole</u> - between two polar molecules (polar=asymmetrical) <u>LDFs</u> - exist in every sample. <u>Bulk Scale Properties</u> - Melting Point, Boiling Point, Viscosity, etc. <u>Types of Solids</u> <u>Kinetic Molecular Theory</u> - gas particles: (1) are far apart (2) are in constant motion (3) collide elastically (4) do not attract or repel each other (5) average k.e. = temperature <u>Ideal Gas Law</u> - $PV = nRT$ <u>Solutions</u> - "like dissolves like" <u>Beer's Law</u> - $A = abc$ represents the change in light's energy as it passes through a material. <u>Photons, wavelength, frequency, and energy</u> - Photons carry energy in waves; $E = hf$ and $c = \lambda \nu$. 	<ul style="list-style-type: none"> <u>Limiting Reactant</u> - compound that runs out during the reaction, stopping it. <u>Writing Net Ionic Equations</u> - Net ionic equations remove spectator ions to show the species that actually interact in a reaction. <u>Combustion Reactions</u> - $\text{Hydrocarbon} + \text{O}_2 \rightarrow \text{H}_2\text{O} + \text{CO}_2$ <u>Redox Reactions</u> - Transfer of electrons. <u>Acid-Base Reactions</u> - Transfer of protons. <u>Precipitation Reactions</u> - Formation of insoluble solids. <u>Stoichiometry</u> - Mole conversions to predict amounts of products or reactants. <u>Titration</u> - Finding an equivalence point for acid-base reactions. 	<ul style="list-style-type: none"> <u>Rates of Reaction</u> - The rate at which reactants turn into products. <u>Rate Laws</u> - Relates to the concentration of reactants and the reaction order. <u>Integrated Rate Laws</u> - Time affects concentration of a reactant. <u>Collision Theory</u> - Particles must collide in the right orientation with enough energy to carry out a reaction. The faster this happens, the faster the reaction rate is. <u>Reaction Mechanisms</u> - Elementary reactions that describe steps in a reaction. <u>Rate Determining Step</u> - The slowest step of the reaction. Limits reaction.
Thermodynamics Unit 6 ↓	Equilibrium Unit 7 ↓	Acids and Bases Unit 8 ↓	Applications of Thermodynamics Unit 9 ↓	Additional Information
<ul style="list-style-type: none"> <u>Specific Heat</u> - energy required to raise the temperature of 1g of a substance by 1°C. <u>Enthalpy of Reaction</u> - ΔH, the amount of heat absorbed or released by a reaction. <u>Calorimetry</u> - Experimental way to measure the enthalpy of reaction ($q = mC\Delta T$) <u>Hess's Law</u> - The total enthalpy of reaction is a sum of the enthalpies for each step. <u>Enthalpy of Formation</u> - The change in enthalpy of forming 1 mole of a compound. <u>Bond Enthalpy</u> = Σ energy of bonds broken - Σ energy of bonds formed 	<ul style="list-style-type: none"> <u>Equilibrium Condition</u> - Forward rate = reverse rate and concentrations are constant. <u>Equilibrium Expression and Constant</u> - Ratio of products to reactants at equilibrium. <u>ICE Tables</u> - Calculate equilibrium concentrations or pressures. <u>Reaction Quotient</u> - Ratio of products to reactants at any point in the reaction. <u>Solubility Product</u> - Ratios/products of soluble compounds. Na, K, NH₄⁺, and nitrate salts are soluble in water. <u>Le Chatelier's Principle</u> - Reactions counteract changes the system in order to maintain equilibrium. 	<ul style="list-style-type: none"> <u>Acids</u> - produce H⁺; H⁺ donors <u>Bases</u> - produce OH⁻; H⁺ acceptors <u>Common Formulas</u> - $\text{pH} = -\log[\text{H}_3\text{O}^+]$, $\text{pOH} = -\log[\text{OH}^-]$, $\text{pH} + \text{pOH} = 14$, $[\text{H}^+][\text{OH}^-] = K_w$. <u>Acid and Base Dissociation Constant</u>: If less than 1, reaction favors the reactants. If greater, favors products. <u>Strong Acids + Bases</u> - completely dissociate into ions in water <u>Percent Dissociation</u> - change in concentration / initial x 100 <u>Buffers</u> - occur between weak substances and their conjugates, they resist drastic changes in pH <u>Henderson-Hasselbalch Equation</u> <u>Titration Curves</u> - pH v volume of titrant added <u>Equivalence Point</u> - $\text{pH} = \text{pK}_a$, $[\text{HA}] = [\text{A}^-]$ 	<ul style="list-style-type: none"> <u>Entropy (ΔS)</u> - disorder The amount of entropy will always increase over time. <u>Gibbs Free Energy (ΔG)</u> - Available energy that can be converted into work Spontaneous = $-\Delta G =$ Thermodynamically favorable $\Delta G = \Delta H - T\Delta S = -RT\ln K$ <u>Voltaic Cells</u> - spontaneous reactions, cell potential must be positive <u>Standard Cell Potential (E°)</u> - potential energy difference between electrodes in volts.. <u>Salt Bridge</u> - balances charge <u>Electrolytic Cells</u> - requires an outside energy source ($I = q/t$). 1 volt = 1 J / 1 coulomb $\Delta G = -nFE^\circ$ 	<p>Content Good to Memorize:</p> <ul style="list-style-type: none"> VSEPR chart Polyatomic ions Equations not on the reference table Strong acids and bases Unit conversions Solubility rules Kinetics - integrated rate laws Kinetics - units of K based on order of reaction Relationship between ΔG, ΔH, and spontaneity <p>AP Format</p> <ul style="list-style-type: none"> MCQ Section - 90 minutes, 60 questions, 50% of the exam FRQ Section - 105 minutes, 7 questions, 50% of the exam, calculator allowed.